DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability and Injury Prevention and Control Special Emphasis Panel: Implementation of the National Occupational Research Agenda (NORA), RFA OH–99–002, Program Area #6 Special Populations at Risk/Aging Workforce

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting.

Name: Disease, Disability and Injury Prevention and Control Special Emphasis Panel: Implementation of the National Occupational Research Agenda (NORA), RFA OH–99–002, Program Area #6 Special Populations at Risk/Aging Workforce.

Times and Dates:

1 p.m.–1:30 p.m., August 4, 1999 (Open)
1:30 p.m.–6 p.m., August 4, 1999 (Closed)

Place: Embassy Suites Hotel, 1900 Diagonal Rd., Alexandria, Va. 22134.

Status: Portions of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to the NORA RFA OH–99–002.

Contact Person For More Information: Michael J. Galvin, Jr., Ph.D., Health Scientist Administrator, Office of Extramural Coordination and Special Projects, NIOSH, CDC, 1600 Clifton Rd., Atlanta, Ga. 30333. Telephone 404/639–3525, e-mail mtg3@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** Notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: July 15, 1999.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–18550 Filed 7–16–99; 1:49 pm]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99F–2336]

Holliday Pigments, Ltd.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Holliday Pigments, Ltd. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of manganese ammonium pyrophosphate (C.I. Pigment Violet 16) as a colorant for all polymers intended for use in contact with food.

DATES: Written comments on the petitioner's environmental assessment by August 20, 1999.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Mark A. Hepp, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3098.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 9B4670) has been filed by Holliday Pigments, Ltd., Morley St., Kingston upon Hull, HU8 8DN ENGLAND. The petition proposes to amend the food additive regulations in §178.3297 *Colorants for polymers* (21 CFR 178.3297) to provide for the safe use of manganese ammonium pyrophosphate (C.I. Pigment Violet 16) as a colorant for all polymers intended for use in contact with food.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before August 20, 1999, submit to the Dockets Management Branch (address above) written comments. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: June 25, 1999.

Alan M. Rulis,

Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 99–18582 Filed 7–20–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99D–1458]

Enforcement Policy: Electronic Records; Electronic Signatures—Compliance Policy Guide; Guidance for FDA Personnel

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a new Compliance Policy Guide (CPG) section 160.850 entitled ''Enforcement Policy: 21 CFR Part 11; Electronic Records; Electronic Signatures.'' This CPG is intended to represent the agency's current thinking on how to comply with the regulations for electronic records and electronic signatures. It also provides that agency decisions on whether or not to pursue regulatory actions will be based on a case-by-case evaluation. The text of the CPG is included in this document.

DATES: Written comments may be submitted at any time.

ADDRESSES: Submit written requests for single copies of CPG section 160.850 entitled ''Enforcement Policy: 21 CFR Part 11; Electronic Records; Electronic Signatures'' to the Division of Compliance Policy (HFC–230), Office of Enforcement, Office of Regulatory Affairs, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20852. Send two self-addressed